

ZHDANOV, Yu.A.; DOROFEYENKO, G.N.; KOROL'CHENKO, G.A.

Gatalyzed acetylation of polyoxy compounds in the presence of magnesium perchlorate. Dokl. AN SSSR 144 no.5:1050-1052 Je (MIRA 15:6) '62.

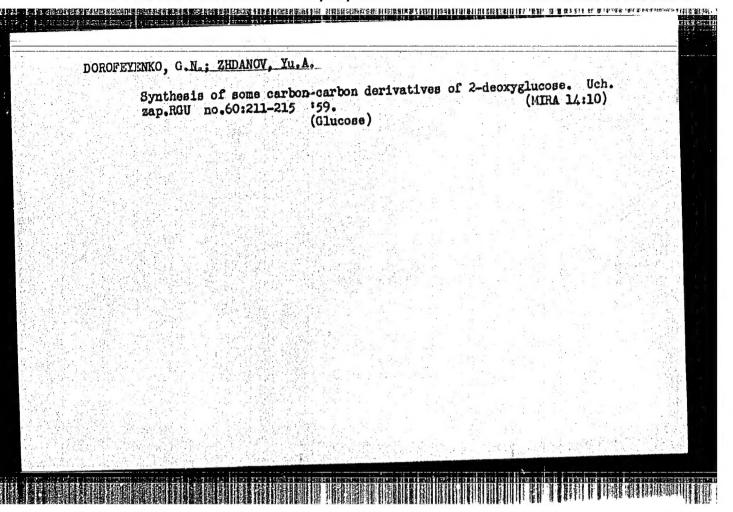
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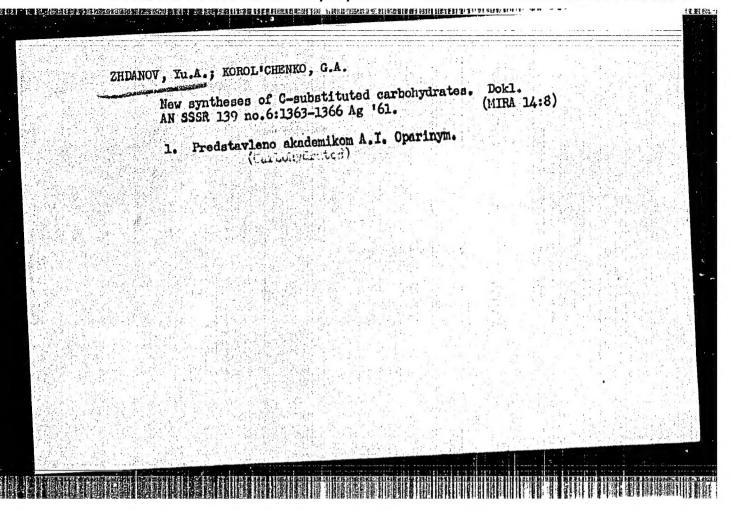
ZHDANOV, Yu.A.; KOROL'CHENKO, G.A.; DOROFEYENKO, C.N.

Gatalytic deacetylation by means of perchloric acid in the carbohydrate series. Dokl. AN SSSR 143 no.4:852-854 Ap (MIRA 15:3) '62.

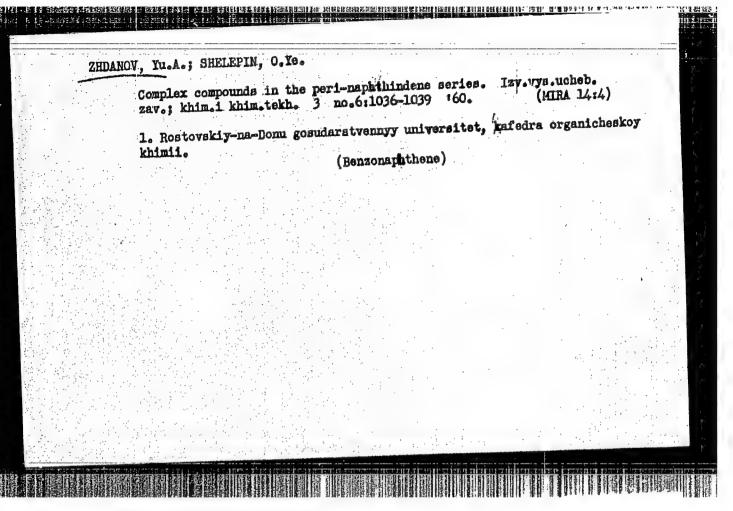
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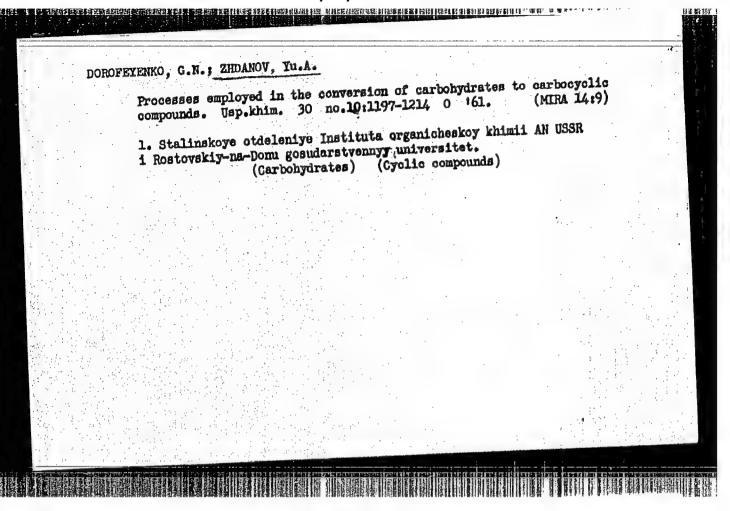
(Acetyl group) (Garbohydrates) (Perchloric acid)

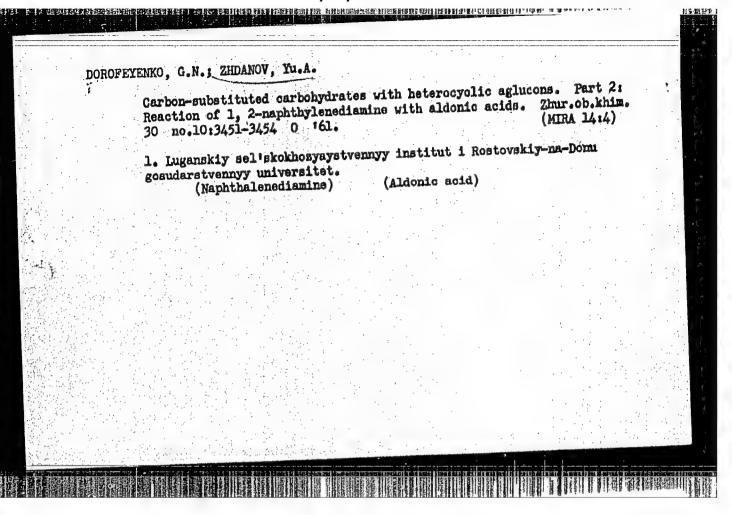




ZHDANOV, Yu. A., (USS	3)				
"Formation of Gly		urbon Atom."			
Report presented at the 10-16 Aug. 1961.	• 5th Int'l. Bi	Lochemistry Co	mgress, Mo	scow,	







# PHASE I BOOK EXPLOITATION

SOV / 5406

# Zhdanov, Yuriy Andreyevich

Ocherki metodologii organicheskoy khimii (Outline Methodology of Organic Chemistry) Moscow, Izd-vo "Vysshaya shkola," 1960. 301 p. Errata slip inserted. 4,700 copies printed.

Ed.: I.K. Korobitsyna; Ed. of Publishing House: A.A. Chiknoverova; Tech, Ed.: I. F. Mulinova.

PURPOSE: This book is intended as supplementary reading matter for chemistry students in schools of higher education.

COVERAGE: The book deals with the methodology of organic chemistry and covers chemical structure, molecular theory, chemical bond, homology, isomerism, and chemical reaction. It presents these concepts from the point of view of Soviet philosophy. The material is based on A. M. Butle-

Card 1/4

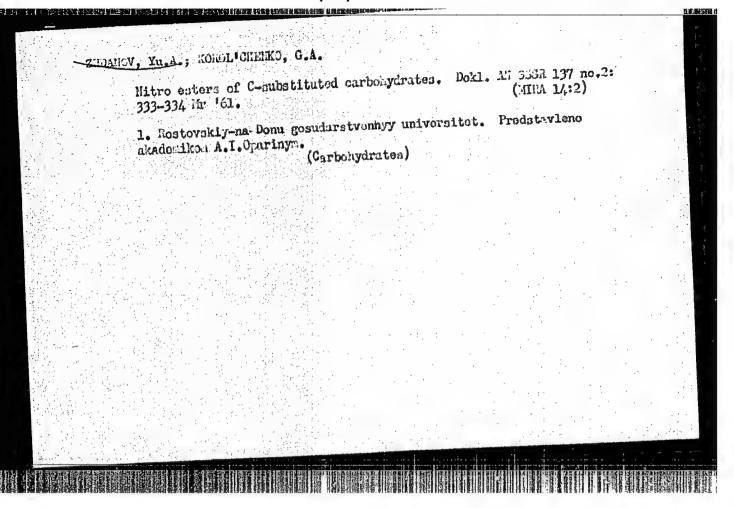
#### Outline Methodology of (Cont.)

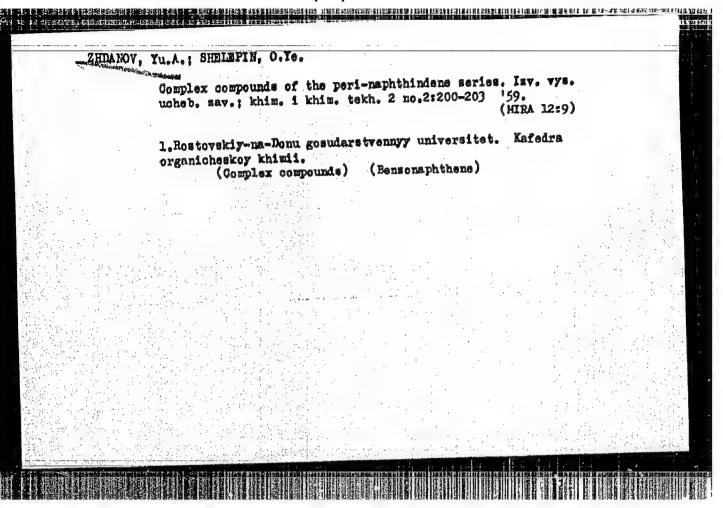
SOV/5406

rov's chemical structure theory which the author considers the foundation of organic chemistry. Much of the material is devoted to reviewing the materialistic and idealistic viewpoints relating to science in general and organic chemistry in particular. No personalities are mentioned. There is no bibliography.

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ZHDANOV, Yu.A.; SHELEPIN, O.Ye.; BAGDASAROV, K.N.; BUDNYATSKAYA, N.I.

Study of the indicator properties of 2-oxy-peri-naphthindenone.
Dokl. AN SSSR 153 no.5:1073-1076 D'63. (MIRA 17:1)

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5(3) AJTHORS:

Zhdanov. Yu. A., Dorofeyenko, G. N.

507/79-29-8-50/81

TITLE:

Some 2,2'-(Polyoxy-alkylene)-dibenzimidazoles

PERIODICAL:

Zhurnal obshchey khimii, 1959, Vol 29, Nr 8, pp 2677-2681 (USSR)

ABSTRACT:

In the present paper, the authors obtained, according to the method described in reference 2, by condensation of o-phenylene diamine with xylo- and 1-arabotrioxy-glutaric acid, the 2,21-(trioxy-propylene)-dibenzimidazoles hitherto not described. They are of crystalline nature and have a very high melting point. The authors improved the methods of synthesizing the 2-(dioxyothylene)-benzimidazole and 2,2'-(dioxy-ethylene)-dibenzimidazole (according to C. S. Hudson and coworkers, Ref 7), in which connection the yields increased by using a mixture of hydrochloric and orthophosphoric acid as condensing agent, and the chlorides, picrates and the diacetyl derivative of the latter, which have hitherto not been described, were obtained. It was found that some acetylated aldonic and saccharic acids could be identified in the form of imidazole derivatives in good yields. On condensation of c-phenylene diamine, a complete separation of the acetyl groups takes place, yielding the same products as in the reaction

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Some 2,2'-(Polyoxy-alkylene)-dibenzimidazoles

507/79-29-8-50/81

of the aldonic and saccharic acids. Thus the tetraacetyl-mucic acid and its acid dichloride produce dibenzimidazole derivatives when heated with 2 moles of o-phenylene diamine in the presence of mineral acid (50-60% yield). On condensation of the Y-lactone of the tetraacetyl-d-galactonic acid with o-phenylene diamine, the 2-[(d-galacto)-pentoxy-amyl]-benzimidazole (61%) was formed. In addition to the main products, 2-methyl-benzimidazcle was obtained in all cases as side product. The resultant benzimidazole products are readily oxidized by potassium permanganate solution, and give, according to the data of C. F. Huebner and coworkers (Ref 8), benzimidazole-2-carboxylic acid in good yield. 2,21-(dioxy-ethylene)-dibenzimidazole yields, on oxidation with sodium periodide, quantitatively 2-formyl-benzimidazole which is transformed by oxidation of alkaline H202-solution into the benzimidazole-2-carboxylic acid. There are 14 references, 2 of which are Soviet.

ASSOCIATION: Rostovskiy-na-Donu gosudarstvennyy universitet (Rostov-na-Donu State University)

SUBMITTED: Card 2/2 May 17, 1958

5'(4) SOV/20-128-4-23/65 Yu. A., Osipov, O. A., Zhdanov. AUTHORS: Shelepin, O. Ye., Kogan, V. A. The Dipole Moments and Structure of Some Derivatives of TITLE: Perinaphthindenone Doklady Akademii nauk SSSR, 1959, Vol 128, Nr 4, pp 719 - 721 PERIODICAL: (USSR) Perinaphthindenone (I) and benzanthrone (IV) having weak or no ABSTRACT: characteristic ketone properties (Refs 1,2) form very solid complex compounds with protonic and aprotonic acids (Refs 2-4). This suggests a considerable polarity of the C = 0 bond. The instability of perinaphthindene and benzanthrene is expressed by their tendency of passing over into a stable oxidized state. The possible existence of a perinaphthindenyl cation, produced recently as a complex salt (Ref 5), had been presumed earlier (Ref 6) although the attempt at producing it had failed. The calculations of the binding energies in the perinaphthindene

Card 1/4

APPROVED FOR RELEASE: 07/19/2001 CIA-RDP86-00513R002064620018-2"

system by the method of molecular orbits showed that a cationoid state with a  $12\pi$ -electron assembly is energetically advantageous for this system. The system is aromatic if it has this assembly (Ref 7). Thus, an intraionic binding character of  ${}^+\text{C} = {}^-\text{O}$ 

The Dipole Moments and Structure of Some Derivatives 50V/20-128-4-23/65 of Perinaphthindenone

can be assumed (according to Ref 8) for the carbonyl compounds of the perinaphthindene series where the negative charge is localized on the oxygen, while the positive one is distributed over the entire carbon system. An extensive analogy of the properties of tropone (II) and perinaphthindenone permits the reduction of its structure to that of perinaphthindenyl oxide (Ia), using also the analogy with tropil oxide (IIa) (see Diagram). For perinaphthindenone, a considerable dipole moment (in the magnitude of 4D) can be expected, all the more so as tropone has a moment between 4.17 and 4.30 D (Ref 9). To clarify this problem, the authors measured the dipole moments of perinaphthindenone and some of its derivatives. Table 1 presents the results showing that the dipole moment in dioxane is reduced by 0.72 D by the introduction of bromine into the nucleus of perinaphthindenone, and in benzanthrone by 1.19 D. The introduction of a benzonal nucleus reduces it by 0.5-0.6 D. On the other hand, the dipole moment increases by the introduction of an oxy group into position 7 of perinaphthindenone (V). An intramolecular cycle with a hydrogen bond is formed. Thus, the negative charge of the carbonyl oxygen is stabilized,

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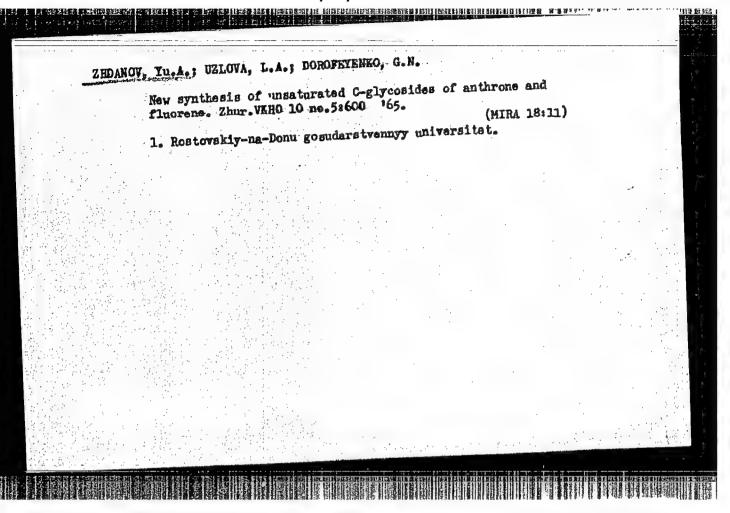
The Dipole Moments and Structure of Some Derivatives SOV/20-128-4-23/65 of Perinaphthindenone

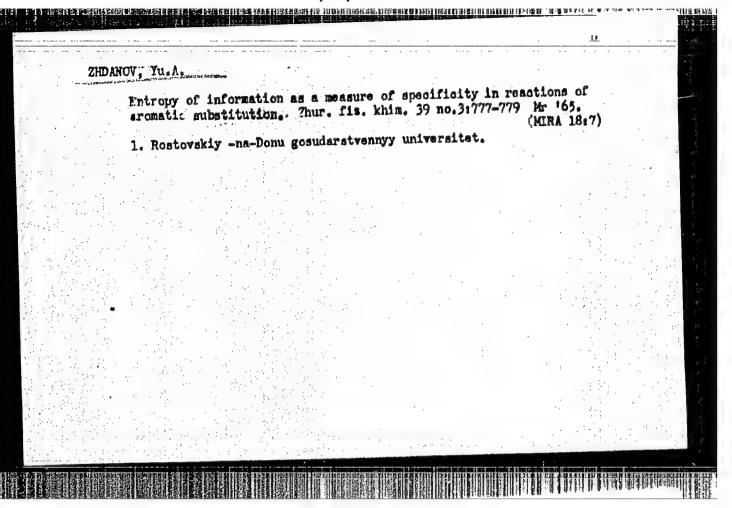
and the C = O group is taken out of the conjugation with the ground skeleton of the molecule due to intracyclic exchange processes via the hydrogen bond. In contrast to the above, the processes via the hydrogen bond. In contrast to the above, the tropolone has a dipole moment much too low (3.7 D) as compared with the tropone. The value of the dipole moment of the complex with the tropone. The value of the dipole moment of the complex with the tropone. The value of the dipole moment of the complex cationoid structure of perinaphthindenylium with a transition of the electron configuration of the antimony atom into the state d<sup>2</sup>sp<sup>3</sup> (similar to HSbCl<sub>6</sub>). The oxygen atom effects a peculiar binding between the cationoid radical of perinaphthindenylium and the antimony atom as one of the addenda of the latter, participating in the coordination sphere with only one of its valences. There are 1 table and 11 references, 6 of which are Soviet.

ASSOCIATION:

Rostovskiy-na-Donu gosudarstvennyy universitet (Rostov-na-Donu State University)

Card 3/4





B/078/60/005/06/24/030 B004/B014

21.3200

Shevchenko, V. B., Shilin, I. V., Zhdanov, Yu. F.

TITLE:

AUTHORS:

The Behavior of Copper Nitrate in the Extraction of the Nitrates of Uranyl and Plutonium by Means of Solutions

of Tributyl Phosphate

PERIODICAL:

Zhurnal neorganicheskoy khimii, 1960, Vol. 5, No. 6,

pp. 1366 - 1374

TEXT: The authors of the present paper wanted to study the behavior of large impurities of copper (in addition to compounds of Ni, Cr, Fe, Co, and Mo) in nuclear fuel that is regenerated by extraction by means of and Mo) in nuclear fuel that is regenerated by extraction by means of benzene—or kerosene solutions of tributyl phosphats (TBP). The authors write down the reaction equation (2) for the extraction of  $\text{Cu}(\text{NO}_3)_2$ , and on the basis of the law of mass action they derive equation (3): and on the basis of the law of mass action they derive equation (5):  $\text{log } K_d = \text{log } K + x \text{ log } \text{TBP} \text{lorg}^3$ , where  $K_d = \text{distribution ratio of Cu}(\text{NO}_3)_2$  and K = equilibrium constant. It follows from Table 1 and Fig. 1 that increases with rising concentration of TBP and increasing ionic Card 1/3

The Behavior of Copper Nitrate in the Extraction \$/078/60/005/06/24/030 of the Nitrates of Uranyl and Plutonium by Means B004/B014 of Solutions of Tributyl Phosphate

strength  $\mu$  of the aqueous solution.  $K_d$  drops, however, with constant  $\mu$ , constant concentration of TBP, and rising concentration of the copper nitrate in the aqueous solution (Figs. 8 and 9).  $K_d$  is higher in TBP-kerosene solution than in TBP benzene (Table 2). It follows from Fig. 2 that by means of TBP benzene copper nitrate is extracted as  $Cu(NO_3)_2$ . JTBP.2H<sub>2</sub>O by  $Cu(NO_3)_2$ . ZTBP.H<sub>2</sub>O, whereas it is extracted as  $Cu(NO_3)_2$ . JTBP.2H<sub>2</sub>O by means of TBP kerosene. These compounds are only stable above -10°C. Fig. 3 shows the effect of HNO<sub>3</sub> on  $K_d$ , Fig. 4 the distribution of HNO<sub>3</sub> among water and TBP in the presence of  $Cu(NO_3)_2$ . Fig. 5 shows that  $K_d$  does not depend on the equilibrium concentration of the H<sup>†</sup> ion. The distribution ratio of copper nitrate is lowered by the presence of uranyl nitrate (Table 3, Fig. 6), whereas aluminum nitrate raises  $K_d$  (Fig. 7). Furthermore, the authors studied the solubility of copper nitrate in TBP as well as the physical data of this solvent (Tables 4-6, Fig. 10). TBP kerosene is divided into two layers when it is saturated

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The Behavior of Copper Nitrate in the Extraction 8/078/60/005/06/24/030 of the Nitrates of Uranyl and Plutonium by Means B004/B014 of Solutions of Tributyl Phosphate

with copper nitrate (Table 7). Hence, the solubility of TBP saturated with copper nitrate is limited in saturated hydrocarbons. There are 10 figures, 7 tables, and 14 references: 8 Soviet, 1 American, 5 British, 1 German, and 1 Yugoslav.

SUEMITTED: February 26, 1959

S/078/60/005/012/014/016 B017/B064

213100 AUTHORS:

Shevchenko, V. B., Shilin, I. V., Zhdanov, Yu. F.

TITLE:

Behavior of Hexavalent and Trivalent Chromium in the Extraction of Uranyl Nitrate and Plutonium Nitrate With Tributyl

Phosphate Solutions

PERIODICAL:

Zhurnal neorganicheskoy khimii, 1960, Vol. 5, No. 12,

pp. 2832-2840

TEXT: Published data (Refs. 2-4), show that in the uranyl nitrate extraction with some organic solvents considerable amounts of chromium are coextracted. The behavior of hexavalent and trivalent chromium in the extraction of uranyl nitrate and plutonium nitrate with tributyl phosphate solutions was studied. The dependence of the distribution coefficient of hexavalent chromium on the tributyl phosphate concentrations was investigated. Hexavalent chromium was found to be extracted with tributyl phosphate, and the distribution coefficient of Cro+ was found to increase when the tributyl phosphate concentration is increased. The effect of concentration of hexavalent chromium upon the distribution coefficient of

Card 1/3

Behavior of Hexavalent and Trivalent Chromium S/078/60/005/012/014/016 in the Extraction of Uranyl Nitrate and B017/B064 Plutonium Nitrate With Tributyl Phosphate Solutions

Cr , and the effect of hydrogen ion concentration upon the distribution coefficient were also studied. Data of Table 3 show that the distribution coefficient rises with increasing concentration of hydrogen ions in the aqueous phase. This proves the fact that the extraction of hexavalent chromium occurs in the form of chromic acid. The following extraction equation is given: H2CrO4 + 3 TBP == H2CrO4.3 TBP. Fig. 4 shows the distribution coefficient of hexavalent chromium as a function of the equilibrium concentration of nitric acid in the aqueous phase. From the course of the curve it may be seen that with increased nitric acid concentration the number of associated chromic acid molecules is also increased. The effect of uranyl nitrate upon the distribution coefficient of hexavalent chromium was investigated. At a concentration of uranyl nitrate higher than 1 mole/1, the distribution coefficient of Cr6+ decreases. The effect of the sodium nitrate concentration upon the Cr6+ distribution coefficient was also studied. The dissociation constants  $K_{3}$  and  $K_{4}$  of the chromic were determined, and the following values acid - tributyl phosphate complex Card 2/3

Behavior of Hexavalent and Trivalent Chromium S/078/60/005/012/014/016 in the Extraction of Uranyl Nitrate and B017/B064 Plutonium Nitrate With Tributyl Phosphate

found::

K<sub>3</sub> = [H<sup>+</sup>] water • [HCro<sub>4</sub>] water = 1.26

[H<sub>2</sub>Cro<sub>4</sub>] water = 1.26

[H<sub>2</sub>Cro<sub>4</sub>·3T] org = 0.535

Table 6 gives the experimental results of the extraction of trivalent chromium. The extraction yield of trivalent chromium with tributyl phosphate is low. The solubility of chromium nitrate in tributyl phosphate is shown in Table 7. The solubility of chromium nitrate in tributyl phosphate rises with increasing tributyl phosphate concentration. There are 5 figures, 7 tables, and 10 references: 4 Soviet, 5 US, and 1 British.

SUBMITTED: August 11, 1959

Card 3/3

SHEVCHENKO, V.B.; ZHDANOV, Yu.F.									
	Extraction phosphate. (Sulfurio	of sulfuric a cid a Radiokhimiia 3 no. acid) (Urany	nd uranyl sulfa 1:7-9 '61. 1 phosphate)	te with tribut (MI) (Butyl phospha	71 W 14:3) te)				
				TRI					

s/186/61/003/006/003/010 E040/E185

Shevchenko, V.B., and Zhdanov, Yu.F.

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AUTHORS : TITLE

Behaviour of plutonium during its extraction with amines from sulphate solutions. I. Extraction of sulphuric acid and tetravalent plutonium sulphate with n-trioctylamine (TOA)

PERIODICAL: Radiokhimiya, v.3, no.6, 1961, 676-684 1;1 mixtures (by volume) of the test solution and trioctylamine solvent (dissolved in carbon tetrachloride) were shaken for 5 minutes, the phases formed were separated by TEXT: centrifuging and the concentrations of sulphuric acid in the aqueous and organic phases were determined volumetrically using phenolphthalein as indicator. Equilibrium concentration of An analysis of the plutonium was determined radiometrically. experimental data obtained for the extraction of H2SO4 with n-trioctylamine showed that, assuming practical insolubility of TOA in the aqueous phase, TOA sulphate and bisulphate are formed in the organic phase at sufficiently high concentrations of

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Behaviour of plutonium during its ... E040/E185

sulphuric acid. Equations are derived for the concentration of both sulphuric acid and TOA bisulphate in the organic phase. The equilibrium constants of the two compounds are, respectively, the equilibrium constants of the two compounds are, respectively, the equilibrium constants of the two compounds are, respectively, the equilibrium coefficient of two phases was found to be independent sulphuric acid between the two phases was found to be independent of the sulphate ion concentration, provided that the second of the sulphuric acid dissociation can be ignored. The stage of sulphuric acid dissociation can be ignored. The stage of sulphuric acid dissociation. Data are also reported increase with rising TOA concentration. Data are also reported increase with rising TOA concentration of sulphuric acid with for the effect of Li<sub>2</sub>SO<sub>4</sub> on the extraction of sulphuric acid with trioctylamine. It was assumed that the extraction process of tetravalent plutonium sulphate with n-trioctylamine can be expressed by the equations:

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Behaviour of plutonium during its .. E040/E185

or 
$$2R_3N_0 + H_B^+ + HSO_{\frac{1}{4}B} + Pu_B^{\frac{1}{4}+} + 2SO_{\frac{1}{4}B}^{2-} \rightleftharpoons (R_3NH)_2Pu(SO_{\frac{1}{4}})_3$$
,  $R_3N_0 + R_3NH_2SO_{\frac{1}{4}} + Pu_B^{\frac{1}{4}+} + 2SO_{\frac{1}{4}B}^{2-} \rightleftharpoons (R_3NH)_2Pu(SO_{\frac{1}{4}})_3$ ,  $H_B^+ + HSO_{\frac{1}{4}B}$ 

The value of the extraction constant, deducted by graphic method, is 1.15 x 10<sup>5</sup>. The distribution of plutonium sulphate between the aqueous and organic phases was found to depend on the relative concentrations of the sulphate and bisulphate of TOA: the distribution coefficient diminishes with an increase in the distribution coefficient of TOA bisulphate. Data tabulated for the effect of sulphuric acid concentration on the distribution coefficient of plutonium sulphate show that the distribution coefficient of plutonium sulphate rises at first with increasing coefficient of plutonium sulphate rises at first with increasing the H2SO4 concentration and then drops. This is thought to be due to an initial inhibition of the hydrolysis of Pu<sup>4+</sup> compounds at sulphuric acid concentrations in aqueous media up to 0.1 M and, at higher H2SO4 concentrations in the aqueous phase, the

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Behaviour of plutonium during its ... \$/186/61/003/006/003/010 E040/E185

concentration of the amine bisulphate in the organic phase tends to rise and lowers the extraction efficiency. Curves plotted for the influence of lithium sulphate on the distribution of tetravalent plutonium between the sulphuric acid and trioctylamine phases show initially a sharp rise and then a progressive reduction of the distribution coefficient of Pu<sup>4+</sup>. There are 4 figures, 4 tables and 22 references: 3 Soviet-bloc, 2 Russian translations of non-Soviet-bloc publications, and 17 non-Soviet-bloc. The four most recent English language references read as follows:

Ref. 10: C.F. Coleman, K.B. Brown, J.G. Moore, K.A. Allen. Paper at the Second Geneva Conference no. 15 (P), 510, 1959.

Ref. 15: D.J. Carswell, J.J. Lawrance,

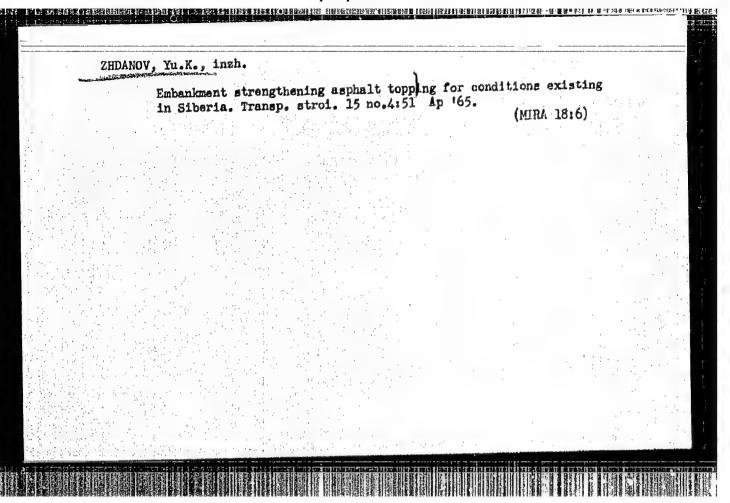
J. Inorg. Nucl. Chem., v. 11, 1, 69 (1959).

Ref. 16; J. L. Drumond, J. Chem. Soc., 3218 (1958).

Ref. 20: D.J. Brown, J. Colloid Sc., v.13, 3, 286 (1958).

SUBMITTED: November 21, 1960

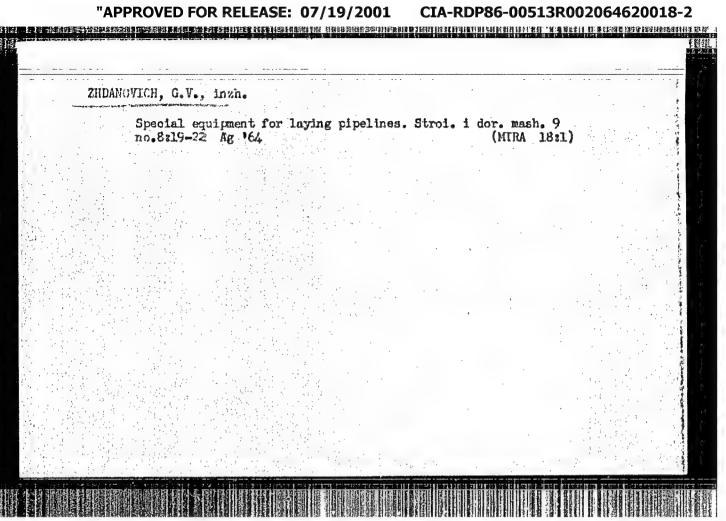
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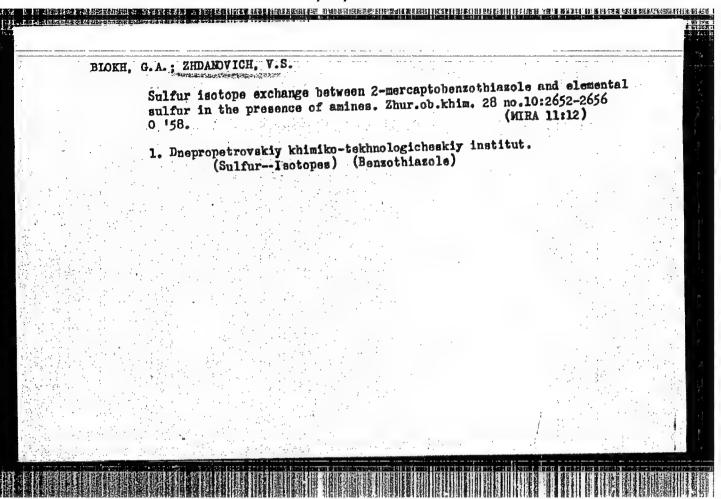


INDOLEV, L.N.; FLEROV, B.L.; ZHDANOV, Yu.Ya.; BROVKIN, A.A.

Herzenbergite from the Deminiskoye deposit. Eckl. AN SSSR 159
no.521044-1047 D '64 (MRA 1821)

1. Yakıtskiy filial Sibirskogo otdeleniya AN SSSR. Fredstavleno akademikem V.I. Smirnovym.





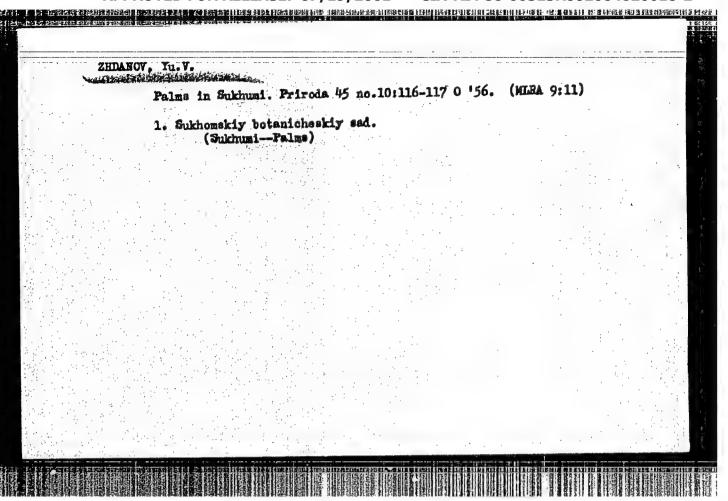
SERGETEV, S. T., kand.tekhn.nauk; ZHADANOV, Yu. S., insh.

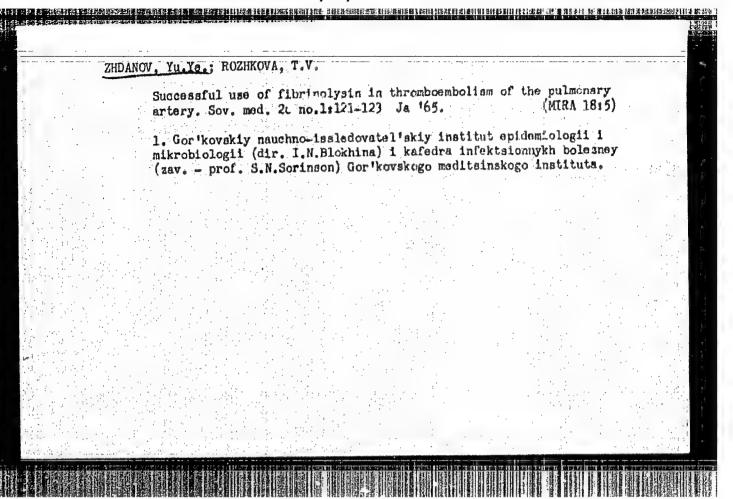
Testing the use of mobile electric substations in the "Proletarskaya-Glubokaya" mine. Ugol' 35 no.7:54-55 Jl '60.

(MIRA 13:7)

1. Shakhta "Proletarskaya-Glubokaya" tresta Makeysvugol'.

(Donets Basin--Electricity in mining) (Electric substations)

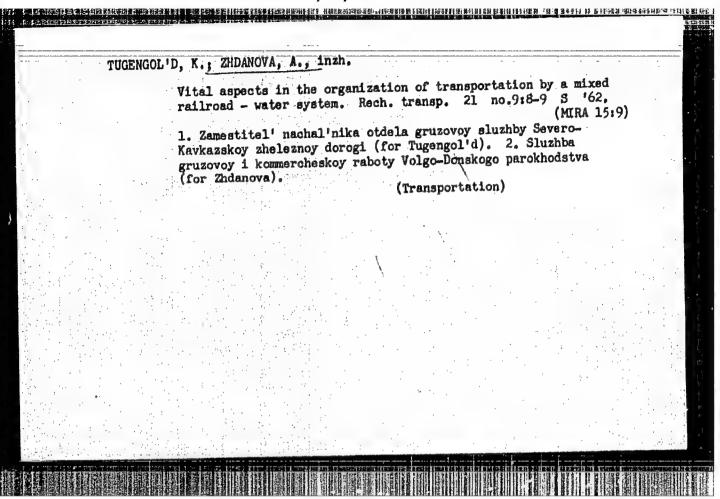




LUKOMSKIY, G.I. (Moskva, K-31, ul. Zhdanova, d.6, kv.6); MANHYICH, A.Z.;
Mikhel'son, V.A.

Hypotonia controlled by arfonad. Nov.khir.arkh. no.5t38-45 8-0 '59.
(MIRA 13:3)

1. Kafedra fakul'tetskop khirurgii (saveduyushchiy - prof. I.8.
Zhorov) sanitarno-giglyenicheskogo fakul'teta l-go Woskovskogo
meditsinskogo instituta.
(HYPOTENSION) (IMIDAZOTHIRMOTHIOLIUM COMPOUNDS)



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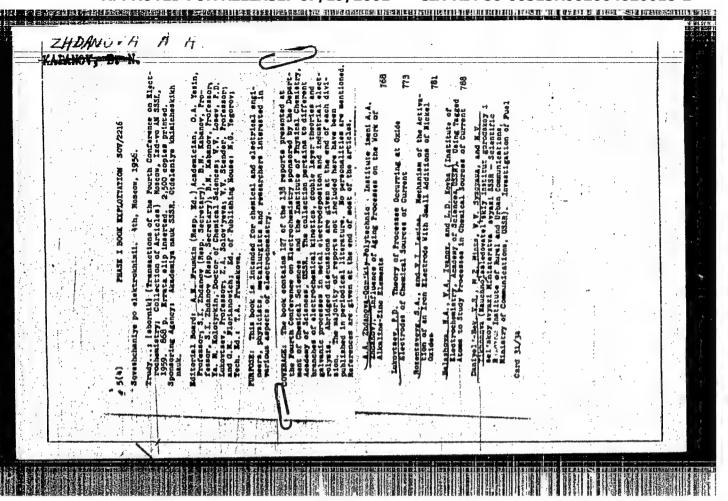
ZHDANOVA, A.A., stershiy nauchnyy sotrudnik; PROKOF'YEV, V.K., prof., doktor fiz.-matem.nauk, otv.red.; FREGER, D.P., tekhn.red.

[Methods of approximate quantitative spectral analysis] Metodika priblizhennogo kolichestvennogo spektral'nogo analiza. Leningrad, 1952. 13 p. (Informatsionno-tekhnicheskii listok, no.72(413)).

(MIRA 14:6)

1. Leningradskiy Dom nauchno-tekhnicheskoy propagandy. 2. Nauchno-issledovatel skiy khimicheskiy institut Leningradskogo gosudarstvennogo universiteta im. A.A.Zhdanova (for Kler). 3. Leningradskiy Dom nauchno-tekhnicheskoy propagandy (for Tyumeneva).

(Spectrum analysis)



MONTHLY List of Russian Accessions Library of Congress October 1952. UNCLASSIFIED

HIDAHOVA, A. D.; NEFEDOVA, T. N.

Textile Industry - Study and Teaching

Instruction in the industrial training school about engineer Kovalev's method. Tekst. prom. 12 No. 7 1952.

Monthly List of Russian Accessions, Library of Congress October 1952, UNCLASSIFIED.

14 16

ZHDANOVA. A. D.; NEFEDOVA, T. N.

Kovalev, Fedor Lukich

Instruction in the industrial training school about engineer Kovalev's method. Tekst. prom. 12, no. 7, 1952.

9. Monthly List of Russian Accessions, Library of Congress, October 1952. 1953, Uncl

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ZHDANOVA, A. D.: NEFEDOVA, T. H.

Textile Industry - Study and Teaching

Instruction in the industrial training school about engineer Kovalev's method. Tedst. prom. 12 No. 7 1952.

9. Monthly List of Russian Accessions, Library of Congress, October 1952 1953, Uncl.

在**经验基础社会转移的现在分词的基础的基础的基础的基础的概念是对于**最后的基础的表现代的。这种的表现的是是一种的是一种的。 M Country : USSR Category: Cultivated Plants. Grains. Abs Jour: RZhBiol., No 22, 1958, No 100261 Zakarpatskaya Oblast State Experimental Agric. Author : Zhdanova, A.F. : A Study of the Size and Shape of the Bed for Corn. Inst (Preliminary Report) Title Orig Pub: Sb. nauchn. tr. Zakarpatsk. obl. gos. s.-kh. opytn. st., 1950-1955 (1957), 1, 18-23 Abstract: The study was conducted in the bottomland zone of Zakarpatakaya Zheltaya Zubovidnaya variety. The best spacing of the plants securing the greatest yield, is the square-pocket of 70 x 70 with two plants to a hill : 1/2 Card M-39

M

Country : USSR

Category: Cultivated Plants. Grains.

Abs Jour: RZhBiol., No 22, 1958, No 100261

and the in-row spacing of 70 x 35 (both variants had 40800 plants on 1 hectare).

: 2/2 Card

### PROVED FOR RELEASE: 07/19/2001 CIA-RDP86-00513R002064620018-2

USSR/Soil Science - Organic Fertilizers.

: Ref Zhur Biol., No 22, 1958, 100097 Abs Jour

: Zhdanova, A.F. Author

Transcarpathian Obl. State Agricultural Experimental Inst Station.

: Utilization of the Green Mass of Lupine in the Capacity of Fertilizers under Tilled Cultivations Title

: Sb. nauchn. tra Zakarpatsk. obl. gos. s.-kh. opytn. st., Orig Pub

1950-1955 (1957), 1, 5-13

Triennial experiments, conducted in the foothill regions of Transcarpathia, indicated that the utilization of the Abstract stubble field's green mass for vegetable fertilization increased the potato crop to 49.9-55.3 c/ha and the corn crop to 8.5-11 c/ha. This increase, at the calculation per unit of fertilization, exceeded considerably

Card 1/2

USSR / Cultivated Plants. Grains.

M-3

Abs Jour: Ref Zhur-Biol., 1958, No 16, 72901.

Author : Zhdanova, A. F. Inst : Zakarpatskaya Oblast State Agricultural Experi-

mental Station.

Title : Periods of Planting Winter Barley.

Orig Pub: Sb. nauchn. tr. Zakarpatsk. obl. gos. s.-kh. opytn. st., 1950-1955 (1957), 1, 13-18.

Abstract: Results of 3 year tests at the station. In the soil-climatic conditions of the lowland rayons of Transcarpathia, the best period for planting winter barley is the period from september 30 to October 10.

Card 1/1

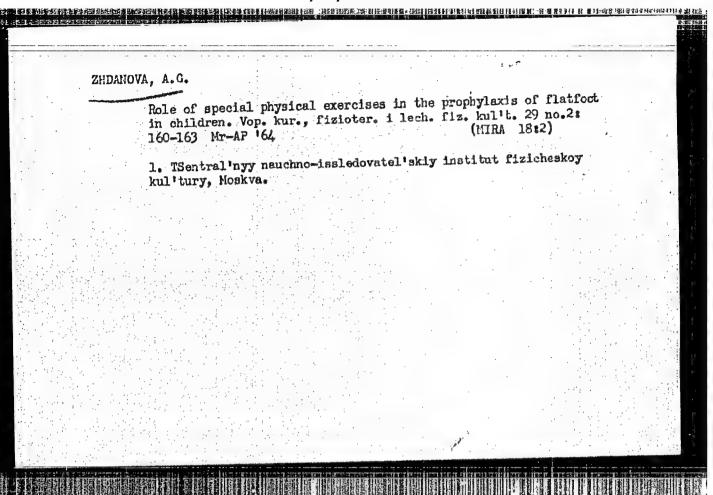
27

ZHDANOVA, A.C. (Moskva, G-146, Komsomol'skiy prospekt, 36, kv.69)

Changes in the body composition of sportswomen due to physical exercise. Arkh. anat., gist. i embr. 46 no.6:113-120 Je '64.

(MIRA 18:3)

1. Gruppa funktsional'noy antropologii (rukovoditel' - kund. biol. nauk A.G. Zhdanova) TSentral'nogo nauchno-issledovatel'-skogo instituta fizicheskoy kul'tury, Moskva.



ZHDANCVA, A. N.

Dissertation: "Concerning the Hydrates of Selenic Acids."

15/12/50

Moscow Order of Lenin Chemicotechnological Inst imeni D. I. Mendeleyev

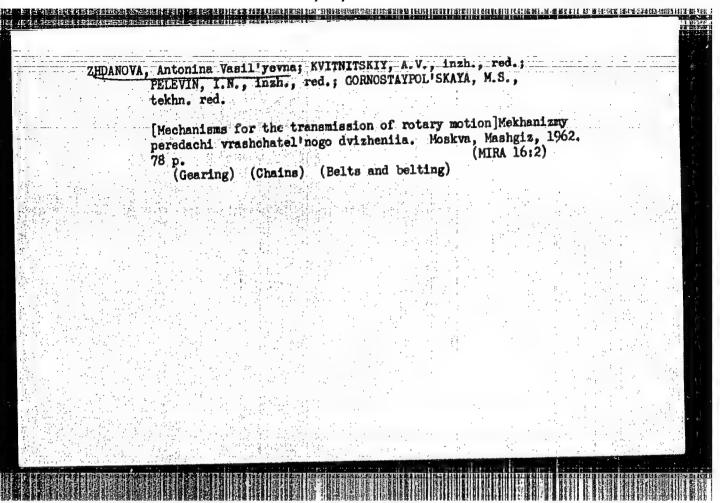
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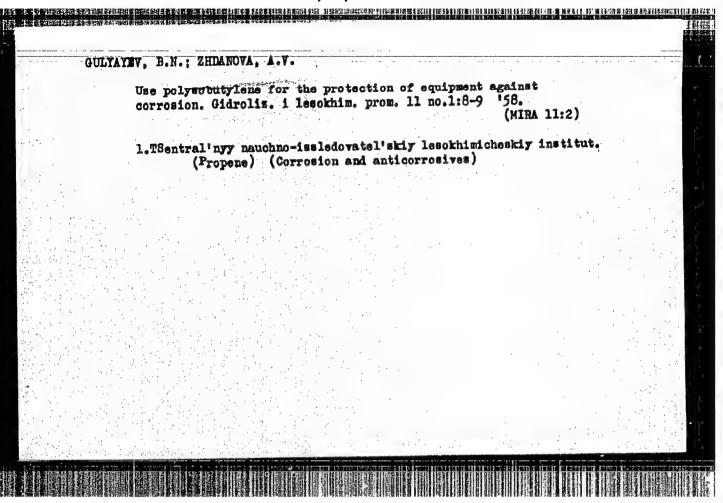
ZHDANOVA, A. H. USSR/Chemistry Oard 1/1 : Zhdanova, A. N. Author : Electrochemical investigation of processes organing on the surface of Title heavy metal-sulfide crystals : Zhur. Fiz. Khim., 28, Ed. 5, 806 - 809, May 1954 Periodical : The changes taking place on the surface of sulfide crystale, during their contact with water containing dissolved gases, were investigated electro-Abstract chemically. It was setablished that the games cause only slight changes in the state of a charita has a second of a charical re-4 - ----44 er thus of wh shar: 15 sulfide surfa e ett to tr 7-USSR: 1-USA. Graphs. Institution : Scientific-Research Institute of Non-Perrous Merallurgy, Moscow : July 13, 1953 Submitted

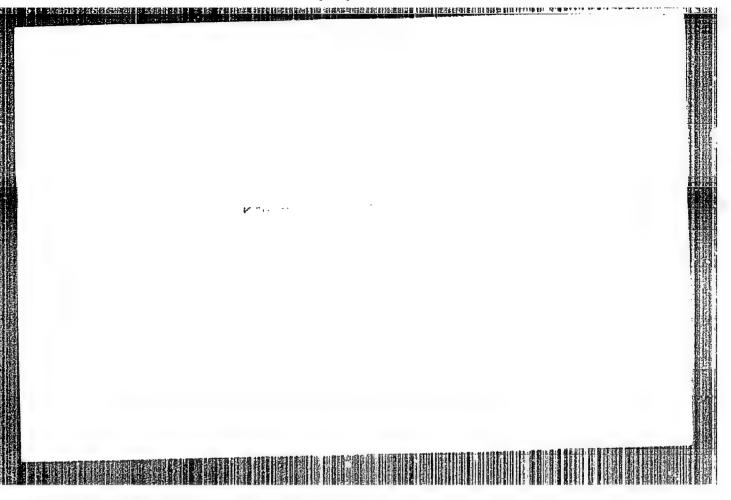
# FEDOROV, Ye.A.; VOLIKOVA, I.G.; ZHDANOVA, A.V. Corrosion resistance of steel materials used in the production of acetic acid. Gidrolis.i lesokhim.prom. 13 no.4:16-19 (MIRA 13:7)

1. Syavskiy lesokhimicheskiy kombinat (for Fedorov). 2. Nauchno-issledovatel'skiy institut khimicheskogo mashinostroyeniya (for Volikova). 3. TSentral'nyy nsuchno-issledovatel'skiy lesokhimicheskiy institut (for Zhdanova).

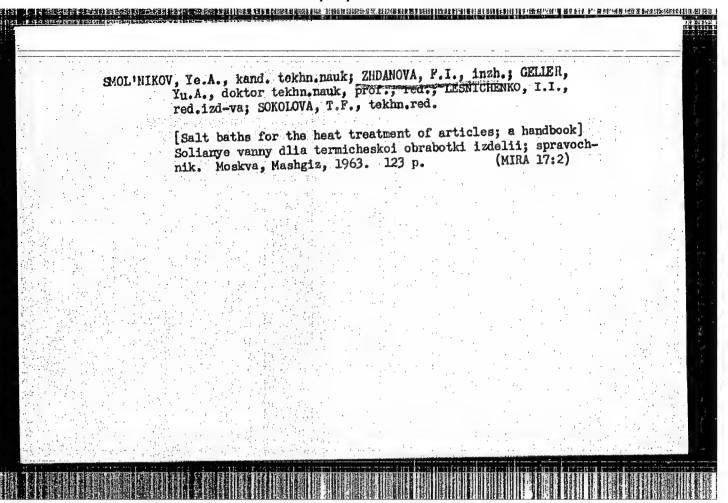
(Steel--Corrosion) (Acetic acid)







ZIDANOVA, B.	
Chemical Abstracts	Study of protein content of awest-cream butter. P. F. D'yuchenko and R. Chilanova. Molochanya Prom. 15. No. 2, 27-6(1964).—In an attempt to insprove the palat-
May 25, 1954 Foods	No. 2, 27-6(1664).—In an attempt to iraprove the palatability and nutritive qualities of sweet-cream butter (I) studies were carried out to det, the effect of manufa, processes on the protein content of L. The results have shown that I made by the continuous-churning or Meleshin's resepa, processes contained more protein than that made in common churas. Owing to I granules picking up coagulated protein in the chura, the continuous churning process had highest protein content of all. D. and Z. also indicated that freshness of I can be detd, by the common of nonprotein and peptide N in the plasma phase. Quant. changes in N substances of the plasma say being studied.  V. N. K.

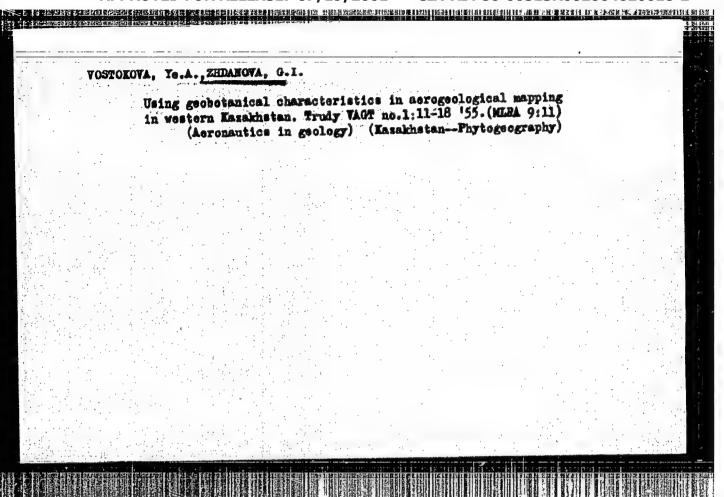


NEYMARK, I.I., prof., red.; KAZANTSEV, I., red.; ZIDANOVA, C., tekhn. red.

[Problems in thoracic and abdominal surgery; collection of works of the Altai Territory Surgical Society] Voprosy grudnoi i briushnoi khirungil; sbornik rabot Altaiskogo kraevogo nauchnogo khirungicheskogo obshchestva. Pod red. I.I.Neimarka. Barnaul, Altaiskoe knizhnoe izd-vo, 1961. 455 p. (MIRA 14:12)

1. Altayskiy kray. Otdel zdravookhraneniya.

(CHEST—SURCERY) (ABDOMEN—SURGERY)



OVCHINNIKOV, Yu.M.; KARPACHEV, S.V.; PAL'GUYEV, S.F.; ZHDANOVA, G.M.; NEUYMIN, A.D.

Kinetics of the reduction by carbon monoxide of solid solutions based on cerium dioxide. Elektrokhimita 1 no.10:1196-1201 0 165.

(MIRA 18:10)

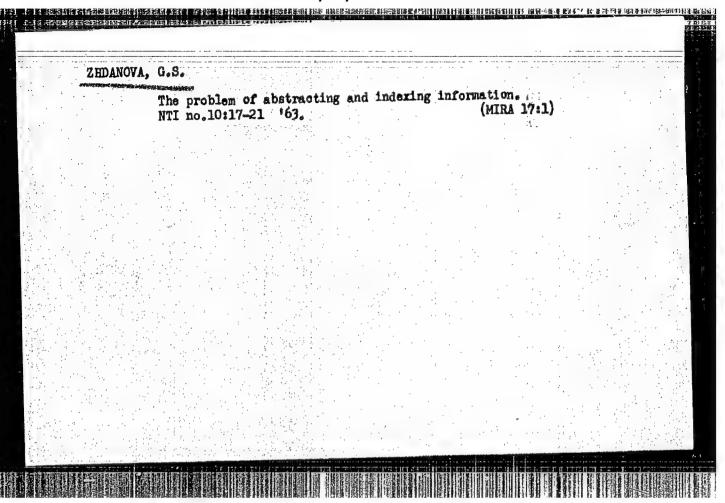
1. Institut elektrokhimii Ural'skogo filiala AN SSSR.

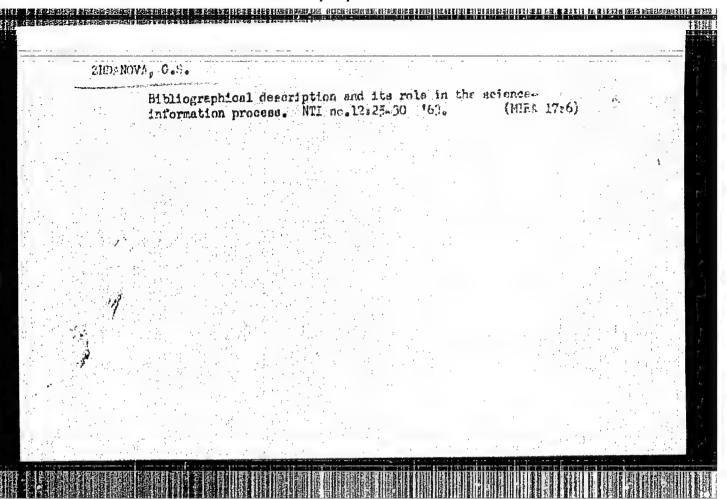
ZHDANOVA, G. P., student IV kursa; KLYASHCHITSKII, A. D., student IV kursa

Device for regulating the screwdriver. Put' i put. khos. 6 (MIRA 15;10)

1. Stroitel'nyy fakul'tet Moskovskogo instituta inzhenerov transporta.

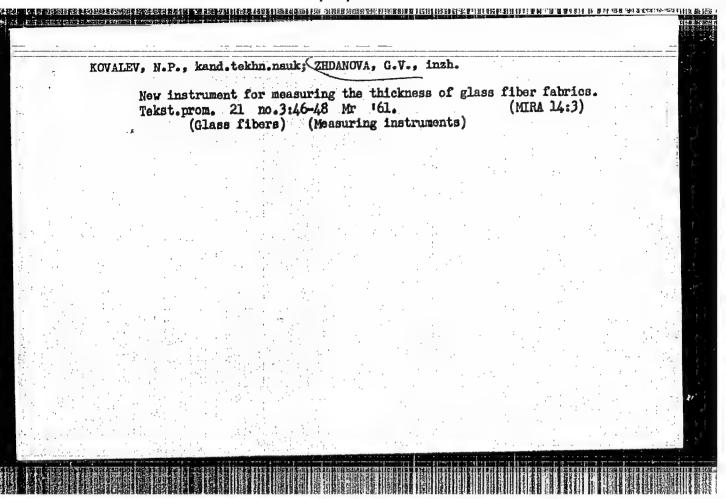
(Railroads—Tools and implements)

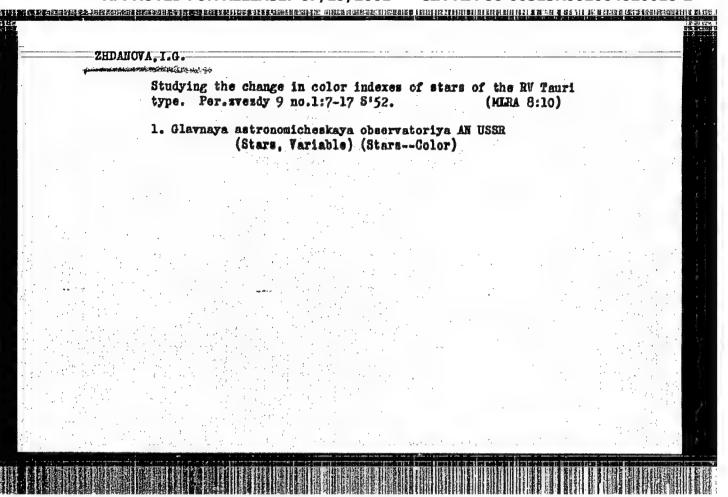


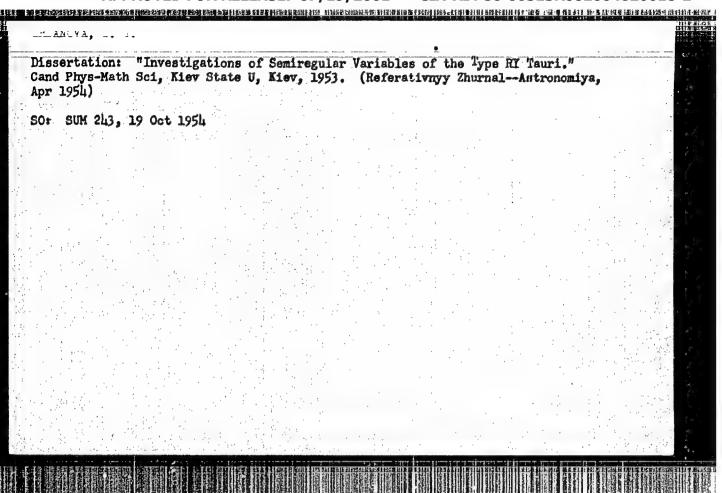


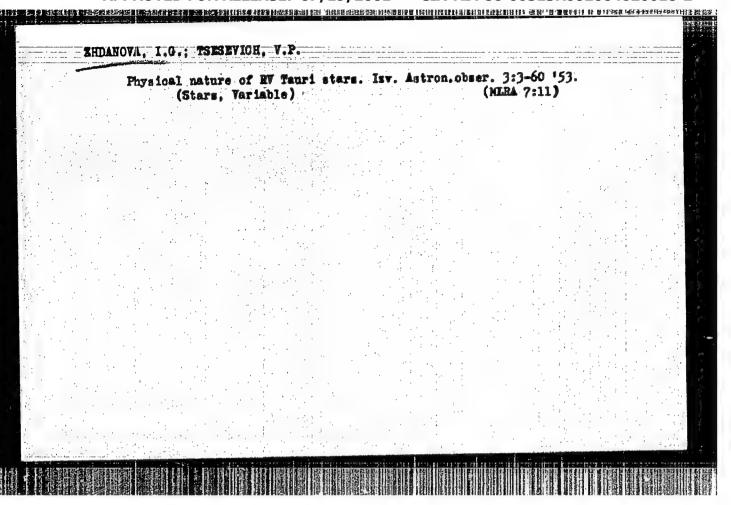
- TUDAHOVA G.S. ISMALZADE. I.G.
- 2. USSR (600)
- h. Silicon Tetraphenyl
- 7. Crystal structure of ortanometallic compounds. Part 2. X-ray investigation of the crystal structure of tetraphenyl compounds of silicon, tin, and lead. Zhur, fiz.khim. 26 no.11,1952.

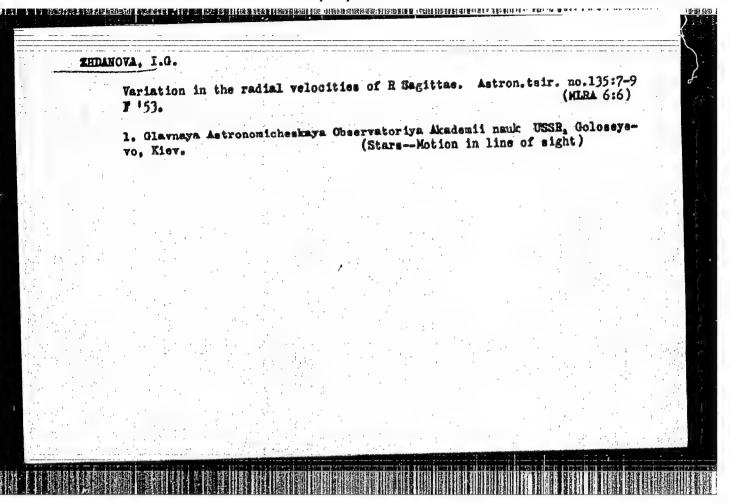
9. Monthly List of Russian Accessions, Library of Congress, April 1953, unclass.

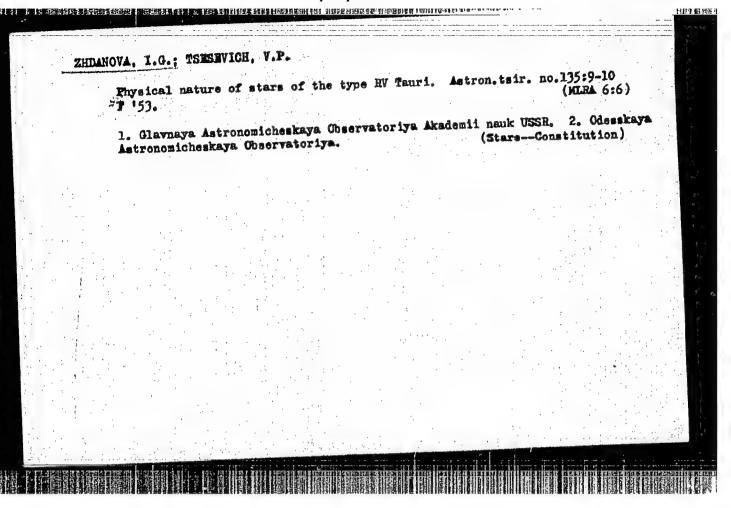


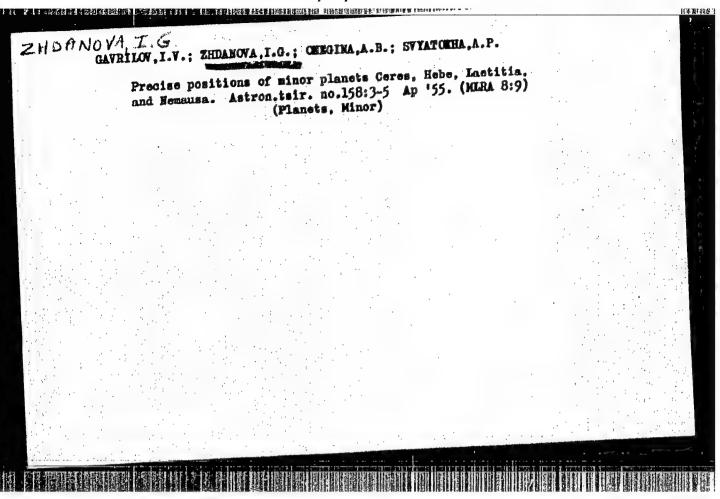


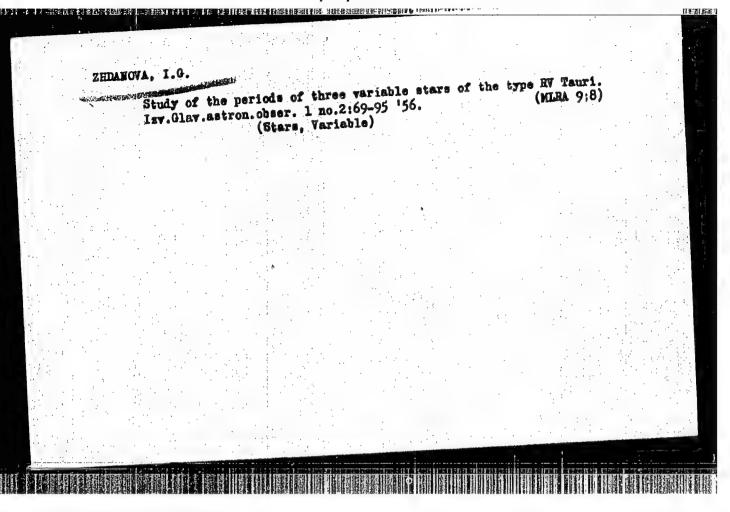












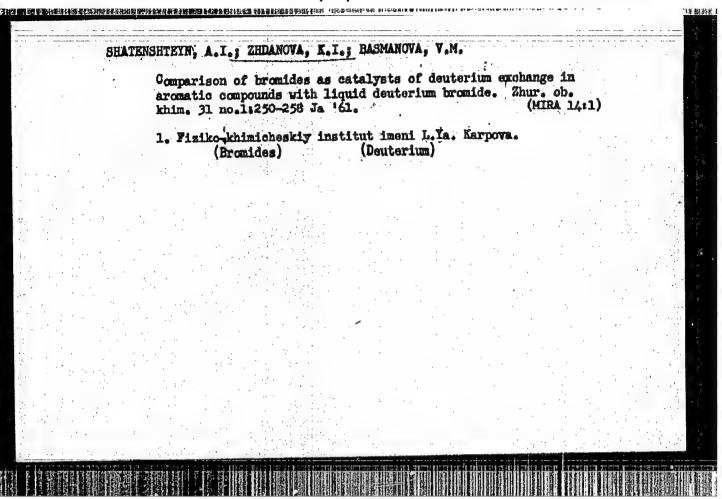
SHATENSHTEYN, A.I., prof.; VYRSKIY, Yu.P., kand. khim. nauk; PRAVIKOVA, N.A., kand. tekhn. nauk; ALIKHANOV, P.F., kand. khim. nauk; ZHDANOVA, K.I., kand. khim. nauk; IZYUMNIKOV, A.L., mlad. nauchn. sotr.; LEVINSKIY, Yu.V., red.

[Practical laboratory manual on the determination of the molecular weights and molecular weight distribution of polymers] Prakticheskoe rukovodstvo po opredeleniiu molekuliarnykh vesov i molekuliarno-vesovogo raspredeleniia polimerov. [By] A.I.Shatenshtein i dr. Moskva, Khimiia, 1964. 188 p. (MIRA 18:2)

ZHDANOVA, K.I.; BASMANOVA, V.M.; SHATSNSHTEYN, A.I.

Catalytic isomerization of methylcyclopentane in liquid hydrogen bromide. Zhur.ob.khim. 31 no.7:2134-2138 Jl '61. (MIRA 14:7)

1. Fiziko-khimicheskiy institut imeni L.Ya. Karpova. (Cyclopentane) (Cyclohexane)



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S/079/61/031/001/022/025 B001/B066

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AUTHORS:

Shatenshteyn, A. I., Zhdanova, K. I., and Basmanova, V. M.

TITLE:

Comparison of Some Bromides as Catalysts in the Deuterium Exchange Between Aromatic Compounds and Liquid Deuterobromide

PERIODICAL:

Zhurnal obshchey khimii, 1961, Vol. 31, No. 1, pp. 250 - 258

TEXT: Only few data are available on the acid catalysis of the isotopic exchange of hydrogen in CH-bonds of organic compounds. The present paper bases upon those by M. Polanyi and co-workers (Ref. 2), by A. Klit, A. Langseth (Ref. 3), and by Shatenshteyn (Ref. 4). The following order of catalytic activity of bromides was established by means of deuterium exchange between liquid deuterobromide and benzene:

AlBr<sub>3</sub>(5 · 10<sup>5</sup>)  $\geqslant$  GaBr<sub>3</sub>(10<sup>5</sup>)  $\geqslant$  FeBr<sub>3</sub>(10<sup>4</sup>)  $\geqslant$  BBr<sub>3</sub>(3 · 10<sup>1</sup>)  $\geqslant$  SbBr<sub>3</sub>(6)  $\geqslant$  TiBr<sub>4</sub>(1)  $\geqslant$  SnBr<sub>4</sub>. The numbers in brackets denote by how many times the deuterium exchange with the given bromide proceeds more quickly than with a TiBr<sub>4</sub> solution of the same concentration. SnBr<sub>4</sub> does not markedly accelerate the reaction.

Card 1/3

-88488

Comparison of Some Bromides as Catalysts in the Deuterium Exchange Between Aromatic Compounds and Liquid Deuterobromide S/079/61/031/001/022/025 B001/B066

InBr, is one of the most active catalysts. The resultant data characterizing the relative electrophilic ratio of the bromides are compared with published data on their relative acidity. The catalysis of hydrogen exchange in aromatic compounds with acid-like bromides dissolved in liquid DBr is explained by the formation of complexes consisting of an aromatic compound, deuterobromide, and bromide. Owing to the coordinated unsaturated state of the bromide, and to the relationship between hydrocarbon and deuteron, the D-Br bond is polarized or split, which favors the passing of deuterium into the aromatic nucleus. The formation of a bond between the functional group of the aromatic compound (C6H5NO2;

C<sub>6</sub>H<sub>5</sub>COOH) and the bromide suppresses the catalytic activity of the latter and retards the hydrogen exchange in the aromatic ring. The data obtained agree with the assumption that one and the same reaction of hydrogen exchange had to proceed according to the associative or to the ionic mechanism, depending on its accomplishment. An overlapping of the mechanisms and the formation of intermediates is possible in this connection.

Card 2/3

88148

Comparison of Some Bromides as Catalysts in the Deuterium Exchange Between Aromatic Compounds and Liquid Deuterobromide 5/079/61/031/001/022/025 B001/B066

P. P. Alikhanov is mentioned. There are 1 figure, 8 tables, and 64 references: 21 Soviet, 24 US, 13 British, 6 German, and 1 French.

ASSOCIATION: Fiziko-khimicheskiy institut imeni L. Ya. Karpova (Physicochemical Institute imeni L. Ya. Karpov)

SUBMITTED: January 29, 1960

Card 3/3

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**建建全部 (15/2) (1** 

Zhdanova, K. L., Basmanova, V. M., Shatonshteyn, A. I. 28(4) AUTHORS: Method of Taking Weighed Samples From Substances Which Easily React With Air Moisture (Sposob vzyatiya navesok veshchestv, TITLE: legko reagiruyushchikh s atmosfernoy vlagoy) PERIODICAL: Zhurnal fizicheskoy khimii, 1959, Vol 33, Nr 6, pp 1438 - 1439 (USSR)

This article describes a device (Fig) which permits precisely weighed samples (from 0.0001 to 2 g) of easily melting sub-ABSTRACT: stances to be taken with the exclusion of moisture and air. The device is to be used for physico-chemical investigations with the aid of substances such as the halides of aluminum, tita nium, tin, and similar elements. In principle, the device is a glass vessel in which - under vacuum and after corresponding heating - a glass ampule with the substance is broken at the moment of melting. The liquid substance enters into small weighed glass ampules (up to 20 pieces) which are closed by melting with the aid of a heated wire. After an accurate description of the device and the working procedure, the authors express their thanks to the glass blower A. A. Orlov. There is

ander and the state of the contractive and the contractive and a state of the contractive action and the contractive action act

Method of Taking Weighed Samples From Substances Which SOV/76-33-6-43/44 Easily React With Air Moisture

1 figure.

ASSOCIATION: Fiziko-khimicheskiy institut im. L. Ya. Karpova, Moskva (Physicochemical Institute imeni L. Ya. Karpov, Moscow)

SUBMITTED: December 20, 1958

Card 2/2

照的技术形列。 斯爾克斯 附近的英語的 自由的性质的 重體 自由的结果的现在分词 经自由的 自由的 自由的 医动物 医多元间 。 以,在 这点的是 这一个一个一个一个 5(4) Vol'pin, M. Ye., Zhdanova, K. I., sov/62-59-4-37/42 AUTHORS: Kursanov, D. N., Setkina, V. N., Shatenshteyn, A. I. On the Interaction of Tropilium Salts With Electrophilic Rea-TITLE: gents (O vzaimodeystvii soley tropiliya s elektrofil'nymi reagentami) Izvestiya Akademii nauk SSSR. Otdeleniye khimicheskikh nauk, PERIODICAL: 1959, Nr 4, pp 754-755 (USSR) This is a brief communication on the investigation of the ABSTRACT: deuterium exchange of tropilium salt in anhydrous DoSOA. It was found that at room temperature the tropilium ion does not take part in the reaction of the deuterium exchange even in the course of 168 hours. Thereafter the deuterium exchange was investigated under aggravated conditions, in liquid DBr in the presence of AlBr3. It was found that tropilium bromide does practically not exchange the deuterium even under aggravated conditions, with AlBr3 excess. (The exchange amounts to no more than 0.9 % in the course of 94 hours). The experiments showed a strong restraint of the electrophilic attack in tropilium salts. In this respect tropilium turned out to

On the Interaction of Tropilium Salts With Electro- SOV/62-59-4-37/42 philic Reagents

be considerably more inactive than benzene and even unsaturated hydrocarbons. The cause of such a difficult course of the electrophilic substitution in the tropilium ion might be that all carbon atoms of the tropilium ring have a positive charge and the system has an electron deficit. This is in accordance with the general conception of the effect of the charge on the deuterium exchange (Ref 5). It can be expected that also other electrophilic reactions will be as little characteristic of the tropilium ion and as difficult as the deuterium exchange. There are 7 references, 4 of which are Soviet.

ASSOCIATION:

Institut elementoorganicheskikh soyedineniy Akademii nauk SSSR (Institute of Elemental-organic Compounds of the Academy of Sciences, USSR). Fiziko-khimicheskiy institut im. Karpova (Physico-chemical Institute imeni Karpov)

SUBMITTED:

September 7, 1958

Card 2/2



1496, 1087, 18.3200

S/081/61/000/013/002/028 B105/B201

AUTHORS:

Yatunova V. A., Zhdanova K. P.

TITLE:

Influence of the chromium content on the rate of hydrogen

· 医克拉克曼氏试验检 计设置对路线的电路计划 (程度电影影響的 经经验的股票)的约11张路 经利益的 1816年 1

diffusion in iron-chromium alloys

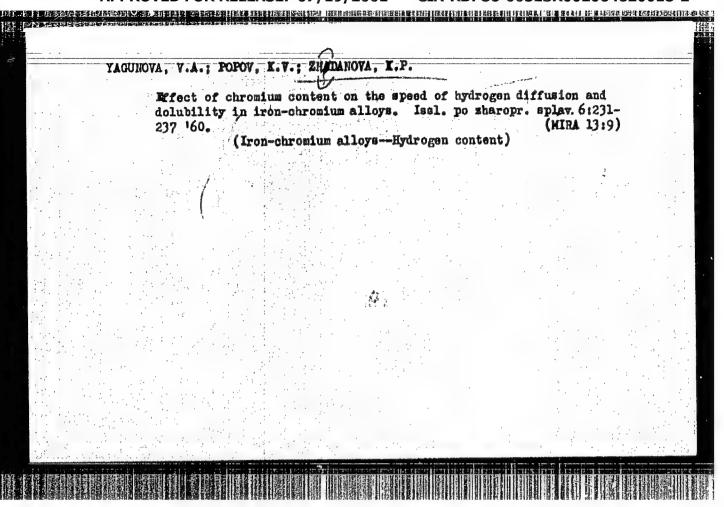
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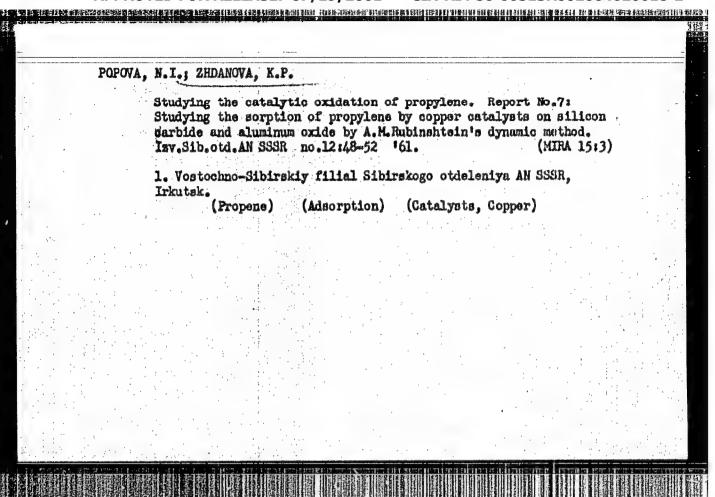
Referativnyy zhurnal. Khimiya, no. 13, 1961, 39, abstract 136259 (Materialy k Konferentsii molodykh nauchn. sotrudn. (Vost.-Sib. fil. Sib. otd. AN SSSR). Vyp. 3. Blagoveshchensk

1960, 89 - 94)

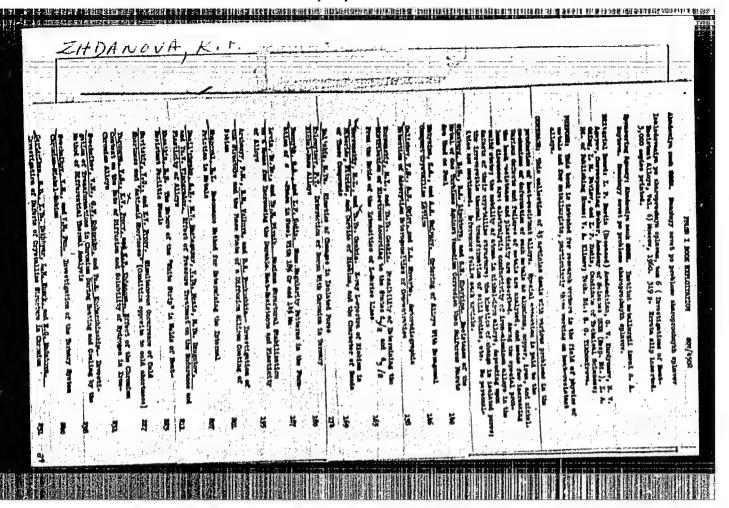
The authors studied the rate of diffusion, D, of hydrogen through this plates of an iron-chromium alloy, and also the influence of the content of Cr (0.5 - 13 %), of the thickness of the plate, of the grain size, and of the surface condition upon D. It was shown that an increase of the content of Cr in alloys lowers D considerably, especially at Cr concentrations up to 1%. The influence of the grain size on D at room temperature is proportional to the extension of the grain boundaries. [Abstracter's note: Complete translation.

Card 1/1

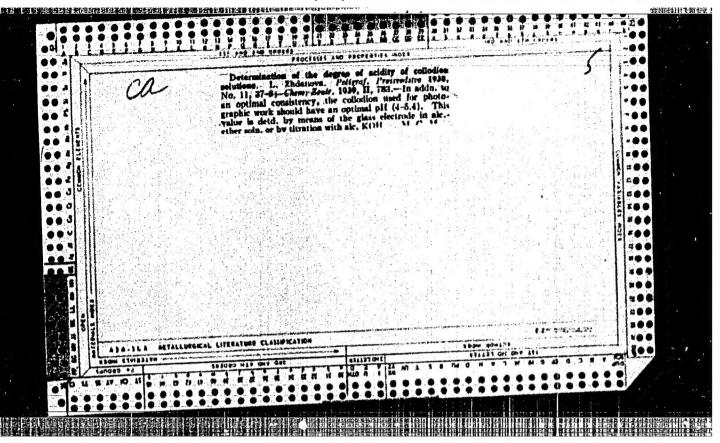




	Studying the crococs of catalytic exidation of propylenc. Report . No.5. Izv.Sib.otd.AN SSSR no.12:72-82 '60. (NIRA 14:2)								
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CHALOV, N.V.; LAPPO-DANILEVSKIY, Yu.K.; GORYACHIKH, Ye.F.; BEINOVA, N.N.; ZHDANOVA, L.A. Chemicomechanical degradation of linters in the presence of sulfuric acid. Sbor. trud. NIIGS 12:87-98 164. (MIRA 18:3)

> CIA-RDP86-00513R002064620018-2" APPROVED FOR RELEASE: 07/19/2001

SHAKH, A.D.; KARASEVA, A.F.; Prinimali uchastiye: ZHDANOVA, L.A.;
NOVOZHILOVA, N.G.; LEBEDEVA, Ye.P.

Technical and economic indices of the rubber goods industry for 1960. Kauch. 1 rez. 20 no.9:41-45 8 161. (MIRA 15:2)

1. Nauchno-issledovatel'skiy institut rezinovoy promyshlennosti.
(Rubber goods)
(Rubber industry-labor productivity)